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US v Navarro et al 20cr160

Motion to Dismiss 1-2

1	Indian Self-Determination and Education Assistance
2	Act (25 U.S.C. 5304).
3	(2) Reclamation state.—The term "Rec-
4	lamation State" means a State or territory described
5	in the first section of the Act of June 17, 1902 (32
6	Stat. 388, chapter 1093; 43 U.S.C. 391).
7	(3) Secretary.—The term "Secretary" means
8	the Secretary of the Interior.
9	TITLE XII—HORSERACING
10	INTEGRITY AND SAFETY
11	SEC. 1201. SHORT TITLE.
12	This title may be cited as the "Horseracing Integrity
13	and Safety Act of 2020".
14	SEC. 1202. DEFINITIONS.
15	In this Act the following definitions apply:
16	(1) Authority.—The term "Authority" means
17	the Horseracing Integrity and Safety Authority des-
18	ignated by section 1203(a).
19	(2) Breeder.—The term "breeder" means a
20	person who is in the business of breeding covered
21	horses.
22	(3) Commission.—The term "Commission"
23	means the Federal Trade Commission.
24	(4) COVERED HORSE.—The term "covered
25	horse" means any Thoroughbred horse, or any other

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1	horse made subject to this Act by election of the ap-
2	plicable State racing commission or the breed gov-
3	erning organization for such horse under section
4	1205(k), during the period—
5	(A) beginning on the date of the horse's
6	first timed and reported workout at a racetrack
7	that participates in covered horseraces or at a
8	training facility; and
9	(B) ending on the date on which the Au-
10	thority receives written notice that the horse
11	has been retired.
12	(5) COVERED HORSERACE.—The term "covered
13	horserace" means any horserace involving covered
14	horses that has a substantial relation to interstate
15	commerce, including any Thoroughbred horserace
16	that is the subject of interstate off-track or advance
17	deposit wagers.
18	(6) COVERED PERSONS.—The term "covered
19	persons" means all trainers, owners, breeders, jock-
20	eys, racetracks, veterinarians, persons (legal and
21	natural) licensed by a State racing commission and
22	the agents, assigns, and employees of such persons
23	and other horse support personnel who are engaged
24	in the care, training, or racing of covered horses.

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1	(7) Equine constituencies.—The term
2	"equine constituencies" means, collectively, owners,
3	breeders, trainers, racetracks, veterinarians, State
4	racing commissions, and jockeys who are engaged in
5	the care, training, or racing of covered horses.
6	(8) Equine industry representative.—The
7	term "equine industry representative" means an or-
8	ganization regularly and significantly engaged in the
9	equine industry, including organizations that rep-
10	resent the interests of, and whose membership con-
11	sists of, owners, breeders, trainers, racetracks, vet-
12	erinarians, State racing commissions, and jockeys.
13	(9) Horseracing anti-doping and medica-
14	TION CONTROL PROGRAM.—The term "horseracing
15	anti-doping and medication control program" means
16	the anti-doping and medication program established
17	under section 1206(a).
18	(10) Immediate family member.—The term
19	"immediate family member" shall include a spouse,
20	domestic partner, mother, father, aunt, uncle, sib-
21	ling, or child.
22	(11) Interstate off-track wager.—The
23	term "interstate off-track wager" has the meaning
24	given such term in section 3 of the Interstate Horse-
25	racing Act of 1978 (15 U.S.C. 3002).

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1	(12) Jockey.—The term "jockey" means a
2	rider or driver of a covered horse in covered
3	horseraces.
4	(13) Owner.—The term "owner" means a per-
5	son who holds an ownership interest in one or more
6	covered horses.
7	(14) Program effective date.—The term
8	"program effective date" means July 1, 2022.
9	(15) RACETRACK.—The term "racetrack"
10	means an organization licensed by a State racing
11	commission to conduct covered horseraces.
12	(16) RACETRACK SAFETY PROGRAM.—The term
13	"racetrack safety program" means the program es-
14	tablished under section 1207(a).
15	(17) STAKES RACE.—The term "stakes race"
16	means any race so designated by the racetrack at
17	which such race is run, including, without limitation,
18	the races comprising the Breeders' Cup World
19	Championships and the races designated as graded
20	stakes by the American Graded Stakes Committee of
21	the Thoroughbred Owners and Breeders Association.
22	(18) State racing commission.—The term
23	"State racing commission" means an entity des-
24	ignated by State law or regulation that has jurisdic-

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1	tion over the conduct of horseracing within the ap-
2	plicable State.
3	(19) Trainer.—The term "trainer" means an
4	individual engaged in the training of covered horses.
5	(20) Training facility.—The term "training
6	facility" means a location that is not a racetrack li-
7	censed by a State racing commission that operates
8	primarily to house covered horses and conduct offi-
9	cial timed workouts.
10	(21) Veterinarian.—The term "veterinarian"
11	means a licensed veterinarian who provides veteri-
12	nary services to covered horses.
13	(22) WORKOUT.—The term "workout" means a
14	timed running of a horse over a predetermined dis-
15	tance not associated with a race or its first quali-
16	fying race, if such race is made subject to this Act
17	by election under section 1205(k) of the horse's
18	breed governing organization or the applicable State
19	racing commission.
20	SEC. 1203. RECOGNITION OF THE HORSERACING INTEG-
21	RITY AND SAFETY AUTHORITY.
22	(a) In General.—The private, independent, self-
23	regulatory, nonprofit corporation, to be known as the
24	"Horseracing Integrity and Safety Authority", is recog-
25	nized for purposes of developing and implementing a

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1	horseracing anti-doping and medication control program
2	and a racetrack safety program for covered horses, cov-
3	ered persons, and covered horseraces.
4	(b) Board of Directors.—
5	(1) Membership.—The Authority shall be gov-
6	erned by a board of directors (in this section re-
7	ferred to as the "Board") comprised of nine mem-
8	bers as follows:
9	(A) Independent members.—Five mem-
10	bers of the Board shall be independent mem-
11	bers selected from outside the equine industry.
12	(B) Industry members.—
13	(i) In general.—Four members of
14	the Board shall be industry members se-
15	lected from among the various equine con-
16	stituencies.
17	(ii) Representation of equine
18	CONSTITUENCIES.—The industry members
19	shall be representative of the various
20	equine constituencies, and shall include not
21	more than one industry member from any
22	one equine constituency.
23	(2) Chair.—The chair of the Board shall be an
24	independent member described in paragraph (1)(A).

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1	(3) BYLAWS.—The Board of the Authority shall
2	be governed by bylaws for the operation of the Au-
3	thority with respect to—
4	(A) the administrative structure and em-
5	ployees of the Authority;
6	(B) the establishment of standing commit-
7	tees;
8	(C) the procedures for filling vacancies on
9	the Board and the standing committees;
10	(D) term limits for members and termi-
11	nation of membership; and
12	(E) any other matter the Board considers
13	necessary.
14	(c) Standing Committees.—
15	(1) Anti-doping and medication control
16	STANDING COMMITTEE.—
17	(A) IN GENERAL.—The Authority shall es-
18	tablish an anti-doping and medication control
19	standing committee, which shall provide advice
20	and guidance to the Board on the development
21	and maintenance of the horseracing anti-doping
22	and medication control program.
23	(B) Membership.—The anti-doping and
24	medication control standing committee shall be
25	comprised of seven members as follows:

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1	(i) Independent members.—A ma-
2	jority of the members shall be independent
3	members selected from outside the equine
4	industry.
5	(ii) Industry members.—A minority
6	of the members shall be industry members
7	selected to represent the various equine
8	constituencies, and shall include not more
9	than one industry member from any one
10	equine constituency.
11	(iii) QUALIFICATION.—A majority of
12	individuals selected to serve on the anti-
13	doping and medication control standing
14	committee shall have significant, recent ex-
15	perience in anti-doping and medication
16	control rules.
17	(C) Chair.—The chair of the anti-doping
18	and medication control standing committee
19	shall be an independent member of the Board
20	described in subsection $(b)(1)(A)$.
21	(2) RACETRACK SAFETY STANDING COM-
22	MITTEE.—
23	(A) IN GENERAL.—The Authority shall es-
24	tablish a racetrack safety standing committee,
25	which shall provide advice and guidance to the

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1	Board on the development and maintenance of
2	the racetrack safety program.
3	(B) Membership.—The racetrack safety
4	standing committee shall be comprised of seven
5	members as follows:
6	(i) Independent members.—A ma-
7	jority of the members shall be independent
8	members selected from outside the equine
9	industry.
10	(ii) Industry members.—A minority
11	of the members shall be industry members
12	selected to represent the various equine
13	constituencies.
14	(C) Chair.—The chair of the racetrack
15	safety standing committee shall be an industry
16	member of the Board described in subsection
17	(b)(1)(B).
18	(d) Nominating Committee.—
19	(1) Membership.—
20	(A) In general.—The nominating com-
21	mittee of the Authority shall be comprised of
22	seven independent members selected from busi-
23	ness, sports, and academia.
24	(B) Initial membership.—The initial
25	nominating committee members shall be set

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1	forth in the governing corporate documents of
2	the Authority.
3	(C) VACANCIES.—After the initial com-
4	mittee members are appointed in accordance
5	with subparagraph (B), vacancies shall be filled
6	by the Board pursuant to rules established by
7	the Authority.
8	(2) Chair.—The chair of the nominating com-
9	mittee shall be selected by the nominating committee
10	from among the members of the nominating com-
11	mittee.
12	(3) Selection of members of the board
13	AND STANDING COMMITTEES.—
14	(A) Initial members.—The nominating
15	committee shall select the initial members of
16	the Board and the standing committees de-
17	scribed in subsection (c).
18	(B) Subsequent members.— The nomi-
19	nating committee shall recommend individuals
20	to fill any vacancy on the Board or on such
21	standing committees.
22	(e) Conflicts of Interest.—To avoid conflicts of
23	interest, the following individuals may not be selected as
24	a member of the Board or as an independent member of
25	a nominating or standing committee under this section:

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1	(1) An individual who has a financial interest
2	in, or provides goods or services to, covered horses.
3	(2) An official or officer—
4	(A) of an equine industry representative;
5	or
6	(B) who serves in a governance or policy-
7	making capacity for an equine industry rep-
8	resentative.
9	(3) An employee of, or an individual who has a
10	business or commercial relationship with, an indi-
11	vidual described in paragraph (1) or (2).
12	(4) An immediate family member of an indi-
13	vidual described in paragraph (1) or (2).
14	(f) Funding.—
15	(1) Initial funding.—
16	(A) In general.—Initial funding to es-
17	tablish the Authority and underwrite its oper-
18	ations before the program effective date shall be
19	provided by loans obtained by the Authority.
20	(B) Borrowing.—The Authority may bor-
21	row funds toward the funding of its operations.
22	(C) Annual calculation of amounts
23	REQUIRED.—
24	(i) IN GENERAL.—Not later than the
25	date that is 90 days before the program ef-

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fective date, and not later than Novem	ıber
2 1 each year thereafter, the Authority si	hall
determine and provide to each State rac	ing
commission the estimated amount requi	ired
from the State—	
(I) to fund the State's prop	or-
tionate share of the horseracing a	nti-
doping and medication control p	oro-
gram and the racetrack safety p	oro-
gram for the next calendar year;	and
(II) to liquidate the State's p	oro-
portionate share of any loan or fu	nd-
ing shortfall in the current calen	dar
year and any previous calendar ye	ear.
(ii) Basis of Calculation.—	The
amounts calculated under clause (i) shall	ll—
(I) be based on—	
3 (aa) the annual budget	of
the Authority for the follow	ving
calendar year, as approved by	the
Board; and	
(bb) the projected amount	t of
covered racing starts for the y	æar
in each State; and	

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1	(II) take into account other
2	sources of Authority revenue.
3	(iii) Requirements regarding
4	BUDGETS OF AUTHORITY.—
5	(I) Initial budget.—The initial
6	budget of the Authority shall require
7	the approval of $\frac{2}{3}$ of the Board.
8	(II) Subsequent budgets.—
9	Any subsequent budget that exceeds
10	the budget of the preceding calendar
11	year by more than 5 percent shall re-
12	quire the approval of 2/3 of the Board.
13	(iv) Rate increases.—
14	(I) In general.—A proposed in-
15	crease in the amount required under
16	this subparagraph shall be reported to
17	the Commission.
18	(II) NOTICE AND COMMENT.—
19	The Commission shall publish in the
20	Federal Register such a proposed in-
21	crease and provide an opportunity for
22	public comment.
23	(2) Assessment and collection of fees by
24	STATES.—

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1	(A) NOTICE OF ELECTION.—Any State
2	racing commission that elects to remit fees pur-
3	suant to this subsection shall notify the Author-
4	ity of such election not later than 60 days be-
5	fore the program effective date.
6	(B) REQUIREMENT TO REMIT FEES.—
7	After a State racing commission makes a notifi-
8	cation under subparagraph (A), the election
9	shall remain in effect and the State racing com-
10	mission shall be required to remit fees pursuant
11	to this subsection according to a schedule estab-
12	lished in rule developed by the Authority and
13	approved by the Commission.
14	(C) WITHDRAWAL OF ELECTION.—A State
15	racing commission may cease remitting fees
16	under this subsection not earlier than one year
17	after notifying the Authority of the intent of
18	the State racing commission to do so.
19	(D) DETERMINATION OF METHODS.—Each
20	State racing commission shall determine, sub-
21	ject to the applicable laws, regulations, and con-
22	tracts of the State, the method by which the
23	requisite amount of fees, such as foal registra-
24	tion fees, sales contributions, starter fees, and

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1	track fees, and other fees on covered persons,
2	shall be allocated, assessed, and collected.
3	(3) Assessment and collection of fees by
4	THE AUTHORITY.—
5	(A) CALCULATION.—If a State racing com-
6	mission does not elect to remit fees pursuant to
7	paragraph (2) or withdraws its election under
8	such paragraph, the Authority shall, not less
9	frequently than monthly, calculate the applica-
10	ble fee per racing start multiplied by the num-
11	ber of racing starts in the State during the pre-
12	ceding month.
13	(B) Allocation.—The Authority shall al-
14	locate equitably the amount calculated under
15	subparagraph (A) collected among covered per-
16	sons involved with covered horseraces pursuant
17	to such rules as the Authority may promulgate.
18	(C) Assessment and collection.—
19	(i) In general.—The Authority shall
20	assess a fee equal to the allocation made
21	under subparagraph (B) and shall collect
22	such fee according to such rules as the Au-
23	thority may promulgate.
24	(ii) Remittance of Fees.—Covered
25	persons described in subparagraph (B)

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1	shall be required to remit such fees to the
2	Authority.
3	(D) Limitation.—A State racing commis-
4	sion that does not elect to remit fees pursuant
5	to paragraph (2) or that withdraws its election
6	under such paragraph shall not impose or col-
7	lect from any person a fee or tax relating to
8	anti-doping and medication control or racetrack
9	safety matters for covered horseraces.
10	(4) Fees and fines imposed
11	by the Authority shall be allocated toward funding
12	of the Authority and its activities.
13	(5) Rule of Construction.—Nothing in this
14	Act shall be construed to require—
15	(A) the appropriation of any amount to the
16	Authority; or
17	(B) the Federal Government to guarantee
18	the debts of the Authority.
19	(g) Quorum.—For all items where Board approval
20	is required, the Authority shall have present a majority
21	of independent members.
22	SEC. 1204. FEDERAL TRADE COMMISSION OVERSIGHT.
23	(a) In General.—The Authority shall submit to the

Commission, in accordance with such rules as the Com-

25 mission may prescribe under section 553 of title 5, United

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1	States Code, any proposed rule, or proposed modification
2	to a rule, of the Authority relating to—
3	(1) the bylaws of the Authority;
4	(2) a list of permitted and prohibited medica-
5	tions, substances, and methods, including allowable
6	limits of permitted medications, substances, and
7	methods;
8	(3) laboratory standards for accreditation and
9	protocols;
10	(4) standards for racing surface quality mainte-
11	nance;
12	(5) racetrack safety standards and protocols;
13	(6) a program for injury and fatality data anal-
14	ysis;
15	(7) a program of research and education on
16	safety, performance, and anti-doping and medication
17	control;
18	(8) a description of safety, performance, and
19	anti-doping and medication control rule violations
20	applicable to covered horses and covered persons;
21	(9) a schedule of civil sanctions for violations;
22	(10) a process or procedures for disciplinary
23	hearings; and
24	(11) a formula or methodology for determining
25	assessments described in section 1203(f)

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1	(b) Publication and Comment.—
2	(1) In General.—The Commission shall—
3	(A) publish in the Federal Register each
4	proposed rule or modification submitted under
5	subsection (a); and
6	(B) provide an opportunity for public com-
7	ment.
8	(2) APPROVAL REQUIRED.—A proposed rule, or
9	a proposed modification to a rule, of the Authority
10	shall not take effect unless the proposed rule or
11	modification has been approved by the Commission.
12	(e) Decision on Proposed Rule or Modifica-
13	TION TO A RULE.—
14	(1) In general.—Not later than 60 days after
15	the date on which a proposed rule or modification is
16	published in the Federal Register, the Commission
17	shall approve or disapprove the proposed rule or
18	modification.
19	(2) Conditions.—The Commission shall ap-
20	prove a proposed rule or modification if the Commis-
21	sion finds that the proposed rule or modification is
22	consistent with—
23	(A) this Act; and
24	(B) applicable rules approved by the Com-
25	mission.

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1	(3) REVISION OF PROPOSED RULE OR MODI-
2	FICATION.—
3	(A) IN GENERAL.—In the case of dis-
4	approval of a proposed rule or modification
5	under this subsection, not later than 30 days
6	after the issuance of the disapproval, the Com-
7	mission shall make recommendations to the Au-
8	thority to modify the proposed rule or modifica-
9	tion.
10	(B) Resubmission.—The Authority may
11	resubmit for approval by the Commission a pro-
12	posed rule or modification that incorporates the
13	modifications recommended under subpara-
14	graph (A).
15	(d) Proposed Standards and Procedures.—
16	(1) In general.—The Authority shall submit
17	to the Commission any proposed rule, standard, or
18	procedure developed by the Authority to carry out
19	the horseracing anti-doping and medication control
20	program or the racetrack safety program.
21	(2) Notice and comment.—The Commission
22	shall publish in the Federal Register any such pro-
23	posed rule, standard, or procedure and provide an
24	opportunity for public comment.

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1	(e) Interim Final Rules.—The Commission may
2	adopt an interim final rule, to take effect immediately,
3	under conditions specified in section 553(b)(B) of title 5,
4	United States Code, if the Commission finds that such a
5	rule is necessary to protect—
6	(1) the health and safety of covered horses; or
7	(2) the integrity of covered horseraces and wa-
8	gering on those horseraces.
9	SEC. 1205. JURISDICTION OF THE COMMISSION AND THE
10	HORSERACING INTEGRITY AND SAFETY AU-
11	THORITY.
12	(a) In General.—Beginning on the program effec-
13	tive date, the Commission, the Authority, and the anti-
14	doping and medication control enforcement agency, each
15	within the scope of their powers and responsibilities under
16	this Act, as limited by subsection (j), shall—
17	(1) implement and enforce the horseracing anti-
18	doping and medication control program and the
19	racetrack safety program;
20	(2) exercise independent and exclusive national
21	authority over—
22	(A) the safety, welfare, and integrity of
2	
23	covered horses, covered persons, and covered

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1	(B) all horseracing safety, performance,
2	and anti-doping and medication control matters
3	for covered horses, covered persons, and covered
4	horseraces; and
5	(3) have safety, performance, and anti-doping
6	and medication control authority over covered per-
7	sons similar to such authority of the State racing
8	commissions before the program effective date.
9	(b) PREEMPTION.—The rules of the Authority pro-
10	mulgated in accordance with this Act shall preempt any
11	provision of State law or regulation with respect to mat-
12	ters within the jurisdiction of the Authority under this
13	Act, as limited by subsection (j). Nothing contained in this
14	Act shall be construed to limit the authority of the Com-
15	mission under any other provision of law.
16	(c) Duties.—
17	(1) In General.—The Authority—
18	(A) shall develop uniform procedures and
19	rules authorizing—
20	(i) access to offices, racetrack facili-
21	ties, other places of business, books,
22	records, and personal property of covered
23	persons that are used in the care, treat-
24	ment, training, and racing of covered
25	horses;

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1	(ii) issuance and enforcement of sub-
2	poenas and subpoenas duces tecum; and
3	(iii) other investigatory powers of the
4	nature and scope exercised by State racing
5	commissions before the program effective
6	date; and
7	(B) with respect to an unfair or deceptive
8	act or practice described in section 1210, may
9	recommend that the Commission commence an
10	enforcement action.
11	(2) Approval of commission.—The proce-
12	dures and rules developed under paragraph $(1)(A)$
13	shall be subject to approval by the Commission in
14	accordance with section 1204.
15	(d) REGISTRATION OF COVERED PERSONS WITH AU-
16	THORITY.—
17	(1) In general.—As a condition of partici-
18	pating in covered races and in the care, ownership,
19	treatment, and training of covered horses, a covered
20	person shall register with the Authority in accord-
21	ance with rules promulgated by the Authority and
22	approved by the Commission in accordance with sec-
23	tion 1204.
24	(2) AGREEMENT WITH RESPECT TO AUTHORITY
25	RULES, STANDARDS, AND PROCEDURES.—Registra-

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1	tion under this subsection shall include an agree-
2	ment by the covered person to be subject to and
3	comply with the rules, standards, and procedures de-
4	veloped and approved under subsection (c).
5	(3) Cooperation.—A covered person reg-
6	istered under this subsection shall, at all times—
7	(A) cooperate with the Commission, the
8	Authority, the anti-doping and medication con-
9	trol enforcement agency, and any respective
10	designee, during any civil investigation; and
11	(B) respond truthfully and completely to
12	the best of the knowledge of the covered person
13	if questioned by the Commission, the Authority,
14	the anti-doping and medication control enforce-
15	ment agency, or any respective designee.
16	(4) Failure to comply.—Any failure of a
17	covered person to comply with this subsection shall
18	be a violation of section 1208(a)(2)(G).
19	(e) Enforcement of Programs.—
20	(1) Anti-doping and medication control
21	ENFORCEMENT AGENCY.—
22	(A) AGREEMENT WITH USADA.—The Au-
23	thority shall seek to enter into an agreement
24	with the United States Anti-Doping Agency
25	under which the Agency acts as the anti-doping

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1	and medication control enforcement agency
2	under this Act for services consistent with the
3	horseracing anti-doping and medication control
4	program.
5	(B) AGREEMENT WITH OTHER ENTITY.—If
6	the Authority and the United States Anti-
7	Doping Agency are unable to enter into the
8	agreement described in subparagraph (A), the
9	Authority shall enter into an agreement with an
10	entity that is nationally recognized as being a
11	medication regulation agency equal in qualifica-
12	tion to the United States Anti-Doping Agency
13	to act as the anti-doping and medication control
14	enforcement agency under this Act for services
15	consistent with the horseracing anti-doping and
16	medication control program.
17	(C) Negotiations.—Any negotiations
18	under this paragraph shall be conducted in
19	good faith and designed to achieve efficient, ef-
20	fective best practices for anti-doping and medi-
21	cation control and enforcement on commercially
22	reasonable terms.
23	(D) Elements of agreement.—Any
24	agreement under this paragraph shall include a
25	description of the scope of work, performance

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1	metrics, reporting obligations, and budgets of
2	the United States Anti-Doping Agency while
3	acting as the anti-doping and medication con-
4	trol enforcement agency under this Act, as well
5	as a provision for the revision of the agreement
6	to increase in the scope of work as provided for
7	in subsection (k), and any other matter the Au-
8	thority considers appropriate.
9	(E) Duties and powers of enforce-
10	MENT AGENCY.—The anti-doping and medica-
11	tion control enforcement agency under an
12	agreement under this paragraph shall—
13	(i) serve as the independent anti-
14	doping and medication control enforcement
15	organization for covered horses, covered
16	persons, and covered horseraces, imple-
17	menting the anti-doping and medication
18	control program on behalf of the Author-
19	ity;
20	(ii) ensure that covered horses and
21	covered persons are deterred from using or
22	administering medications, substances, and
23	methods in violation of the rules estab-
24	lished in accordance with this Act;

1	(iii) implement anti-doping education,
2	research, testing, compliance and adjudica-
3	tion programs designed to prevent covered
4	persons and covered horses from using or
5	administering medications, substances, and
6	methods in violation of the rules estab-
7	lished in accordance with this Act;
8	(iv) exercise the powers specified in
9	section 1206(e)(4) in accordance with that
10	section; and
11	(v) implement and undertake any
12	other responsibilities specified in the agree-
13	ment.
14	(F) TERM AND EXTENSION.—
15	(i) TERM OF INITIAL AGREEMENT.—
16	The initial agreement entered into by the
17	Authority under this paragraph shall be in
18	effect for the 5-year period beginning on
19	the program effective date.
20	(ii) Extension.—At the end of the 5-
21	year period described in clause (i), the Au-
22	thority may—
23	(I) extend the term of the initial
24	agreement under this paragraph for
25	such additional term as is provided by

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1	the rules of the Authority and con-
2	sistent with this Act; or
3	(II) enter into an agreement
4	meeting the requirements of this para-
5	graph with an entity described by sub-
6	paragraph (B) for such term as is
7	provided by such rules and consistent
8	with this Act.
9	(2) Agreements for enforcement by
10	STATE RACING COMMISSIONS.—
11	(A) STATE RACING COMMISSIONS.—
12	(i) Racetrack safety program.—
13	The Authority may enter into agreements
14	with State racing commissions for services
15	consistent with the enforcement of the
16	racetrack safety program.
17	(ii) Anti-doping and medication
18	CONTROL PROGRAM.—The anti-doping and
19	medication control enforcement agency
20	may enter into agreements with State rac-
21	ing commissions for services consistent
22	with the enforcement of the anti-doping
23	and medication control program.
24	(B) Elements of agreements.—Any
25	agreement under this paragraph shall include a

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1	description of the scope of work, performance
2	metrics, reporting obligations, budgets, and any
3	other matter the Authority considers appro-
4	priate.
5	(3) Enforcement of standards.—The Au-
6	thority may coordinate with State racing commis-
7	sions and other State regulatory agencies to monitor
8	and enforce racetrack compliance with the standards
9	developed under paragraphs (1) and (2) of section
10	1207(e).
11	(f) Procedures With Respect to Rules of Au-
12	THORITY.—
13	(1) Anti-doping and medication con-
14	TROL.—
15	(A) In general.—Recommendations for
16	rules regarding anti-doping and medication con-
17	trol shall be developed in accordance with sec-
18	tion 1206.
19	(B) Consultation.—The anti-doping and
20	medication control enforcement agency shall
21	consult with the anti-doping and medication
22	control standing committee and the Board of
23	the Authority on all anti-doping and medication
24	control rules of the Authority.

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1	(2) RACETRACK SAFETY.—Recommendations
2	for rules regarding racetrack safety shall be devel-
3	oped by the racetrack safety standing committee of
4	the Authority.
5	(g) Issuance of Guidance.—
6	(1) The Authority may issue guidance that—
7	(A) sets forth—
8	(i) an interpretation of an existing
9	rule, standard, or procedure of the Author-
10	ity; or
11	(ii) a policy or practice with respect to
12	the administration or enforcement of such
13	an existing rule, standard, or procedure;
14	and
15	(B) relates solely to—
16	(i) the administration of the Author-
17	ity; or
18	(ii) any other matter, as specified by
19	the Commission, by rule, consistent with
20	the public interest and the purposes of this
21	subsection.
22	(2) Submittal to commission.—The Author-
23	ity shall submit to the Commission any guidance
24	issued under paragraph (1).

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1	(3) Immediate effect.—Guidance issued
2	under paragraph (1) shall take effect on the date on
3	which the guidance is submitted to the Commission
4	under paragraph (2).
5	(h) Subpoena and Investigatory Authority.—
6	The Authority shall have subpoena and investigatory au-
7	thority with respect to civil violations committed under its
8	jurisdiction.
9	(i) CIVIL PENALTIES.—The Authority shall develop
10	a list of civil penalties with respect to the enforcement of
11	rules for covered persons and covered horseraces under its
12	jurisdiction.
13	(j) CIVIL ACTIONS.—
14	(1) In general.—In addition to civil sanctions
15	imposed under section 1208, the Authority may
16	commence a civil action against a covered person or
17	racetrack that has engaged, is engaged, or is about
18	to engage, in acts or practices constituting a viola-
19	tion of this Act or any rule established under this
20	Act in the proper district court of the United States,
21	the United States District Court for the District of
22	Columbia, or the United States courts of any terri-
23	tory or other place subject to the jurisdiction of the
24	United States, to enjoin such acts or practices, to
25	enforce any civil sanctions imposed under that sec-

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1	tion, and for all other relief to which the Authority
2	may be entitled.
3	(2) Injunctions and restraining orders.—
4	With respect to a civil action commenced under
5	paragraph (1), upon a proper showing, a permanent
6	or temporary injunction or restraining order shall be
7	granted without bond.
8	(k) Limitations on Authority.—
9	(1) Prospective application.—The jurisdic-
10	tion and authority of the Authority and the Commis-
11	sion with respect to the horseracing anti-doping and
12	medication control program and the racetrack safety
13	program shall be prospective only.
14	(2) Previous matters.—
15	(A) IN GENERAL.—The Authority and the
16	Commission may not investigate, prosecute, ad-
17	judicate, or penalize conduct in violation of the
18	horseracing anti-doping and medication control
19	program and the racetrack safety program that
20	occurs before the program effective date.
21	(B) STATE RACING COMMISSION.—With re-
22	spect to conduct described in subparagraph (A),
23	the applicable State racing commission shall re-
24	tain authority until the final resolution of the
25	matter.

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1	(3) OTHER LAWS UNAFFECTED.—This Act
2	shall not be construed to modify, impair or restrict
3	the operation of the general laws or regulations, as
4	may be amended from time to time, of the United
5	States, the States and their political subdivisions re-
6	lating to criminal conduct, cruelty to animals, mat-
7	ters unrelated to antidoping, medication control and
8	racetrack and racing safety of covered horses and
9	covered races, and the use of medication in human
10	participants in covered races.
11	(l) Election for Other Breed Coverage Under
12	Act.—
13	(1) In general.—A State racing commission
14	or a breed governing organization for a breed of
15	horses other than Thoroughbred horses may elect to
16	have such breed be covered by this Act by the filing
17	of a designated election form and subsequent ap-
18	proval by the Authority. A State racing commission
19	may elect to have a breed covered by this Act for the
20	applicable State only.
21	(2) Election conditional on funding
22	MECHANISM.—A commission or organization may
23	not make an election under paragraph (1) unless the
24	commission or organization has in place a mecha-
25	nism to provide sufficient funds to cover the costs of

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1	the administration of this Act with respect to the
2	horses that will be covered by this Act as a result
3	of the election.
4	(3) APPORTIONMENT.—The Authority shall ap-
5	portion costs described in paragraph (2) in connec-
6	tion with an election under paragraph (1) fairly
7	among all impacted segments of the horseracing in-
8	dustry, subject to approval by the Commission in ac-
9	cordance with section 1204. Such apportionment
10	may not provide for the allocation of costs or funds
11	among breeds of horses.
12	SEC. 1206. HORSERACING ANTI-DOPING AND MEDICATION
13	CONTROL PROGRAM.
	control program. (a) Program Required.—
13	
13 14	(a) Program Required.—
13 14 15	(a) Program Required.— (1) In general.—Not later than the program
13 14 15 16	(a) Program Required.—(1) In general.—Not later than the program effective date, and after notice and an opportunity
13 14 15 16 17	 (a) Program Required.— (1) In General.—Not later than the program effective date, and after notice and an opportunity for public comment in accordance with section 1204,
13 14 15 16 17 18	(a) Program Required.— (1) In General.—Not later than the program effective date, and after notice and an opportunity for public comment in accordance with section 1204, the Authority shall establish a horseracing anti-
13 14 15 16 17 18	(a) Program Required.— (1) In General.—Not later than the program effective date, and after notice and an opportunity for public comment in accordance with section 1204, the Authority shall establish a horseracing anti-doping and medication control program applicable to
13 14 15 16 17 18 19 20	(a) Program Required.— (1) In General.—Not later than the program effective date, and after notice and an opportunity for public comment in accordance with section 1204, the Authority shall establish a horseracing anti-doping and medication control program applicable to all covered horses, covered persons, and covered
13 14 15 16 17 18 19 20 21	(a) Program Required.— (1) In General.—Not later than the program effective date, and after notice and an opportunity for public comment in accordance with section 1204, the Authority shall establish a horseracing anti-doping and medication control program applicable to all covered horses, covered persons, and covered horseraces in accordance with the registration of
13 14 15 16 17 18 19 20 21 22	(a) Program Required.— (1) In General.—Not later than the program effective date, and after notice and an opportunity for public comment in accordance with section 1204, the Authority shall establish a horseracing anti-doping and medication control program applicable to all covered horses, covered persons, and covered horseraces in accordance with the registration of covered persons under section 1205(d).

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1	that is made subject to this Act by election of a
2	State racing commission or the breed governing or-
3	ganization for such horse under section 1205(k), the
4	Authority shall consider the unique characteristics of
5	such breed.
6	(b) Considerations in Development of Pro-
7	GRAM.—In developing the horseracing anti-doping and
8	medication control program, the Authority shall take into
9	consideration the following:
10	(1) Covered horses should compete only when
11	they are free from the influence of medications,
12	other foreign substances, and methods that affect
13	their performance.
14	(2) Covered horses that are injured or unsound
15	should not train or participate in covered races, and
16	the use of medications, other foreign substances, and
17	treatment methods that mask or deaden pain in
18	order to allow injured or unsound horses to train or
19	race should be prohibited.
20	(3) Rules, standards, procedures, and protocols
21	regulating medication and treatment methods for
22	covered horses and covered races should be uniform
23	and uniformly administered nationally.
24	(4) To the extent consistent with this Act, con-
25	sideration should be given to international anti-

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1	doping and medication control standards of the
2	International Federation of Horseracing Authorities
3	and the Principles of Veterinary Medical Ethics of
4	the American Veterinary Medical Association.
5	(5) The administration of medications and
6	treatment methods to covered horses should be
7	based upon an examination and diagnosis that iden-
8	tifies an issue requiring treatment for which the
9	medication or method represents an appropriate
10	component of treatment.
11	(6) The amount of the apeutic medication that
12	a covered horse receives should be the minimum nec-
13	essary to address the diagnosed health concerns
14	identified during the examination and diagnostic
15	process.
16	(7) The welfare of covered horses, the integrity
17	of the sport, and the confidence of the betting public
18	require full disclosure to regulatory authorities re-
19	garding the administration of medications and treat-
20	ments to covered horses.
21	(c) ACTIVITIES.—The following activities shall be car-
22	ried out under the horseracing anti-doping and medication
23	control program:
24	(1) STANDARDS FOR ANTI-DOPING AND MEDI-
25	CATION CONTROL.—Not later than 120 days before

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1	the program effective date, the Authority shall issue,
2	by rule—
3	(A) uniform standards for—
4	(i) the administration of medication to
5	covered horses by covered persons; and
6	(ii) laboratory testing accreditation
7	and protocols; and
8	(B) a list of permitted and prohibited
9	medications, substances, and methods, including
10	allowable limits of permitted medications, sub-
11	stances, and methods.
12	(2) Review process for administration of
13	MEDICATION.—The development of a review process
14	for the administration of any medication to a cov-
15	ered horse during the 48-hour period preceding the
16	next racing start of the covered horse.
17	(3) AGREEMENT REQUIREMENTS.—The devel-
18	opment of requirements with respect to agreements
19	under section 1205(e).
20	(4) Anti-doping and medication control
21	ENFORCEMENT AGENCY.—
22	(A) CONTROL RULES, PROTOCOLS, ETC.—
23	Except as provided in paragraph (5), the anti-
24	doping and medication control program enforce-
25	ment agency under section 1205(e) shall, in

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3017 1 consultation with the anti-doping and medica-2 tion control standing committee of the Author-3 ity and consistent with international best prac-4 tices, develop and recommend anti-doping and 5 medication control rules, protocols, policies, and 6 guidelines for approval by the Authority. 7 (B) RESULTS MANAGEMENT.—The anti-8 doping and medication control enforcement 9 agency shall conduct and oversee anti-doping 10 and medication control results management, in-11 cluding independent investigations, charging 12 and adjudication of potential medication control 13 rule violations, and the enforcement of any civil 14 sanctions for such violations. Any final decision 15 or civil sanction of the anti-doping and medication control enforcement agency under this sub-16 17 paragraph shall be the final decision or civil 18 sanction of the Authority, subject to review in 19 accordance with section 1209. 20 (C) Testing.—The anti-doping enforce-21 ment agency shall perform and manage test dis-22

(C) Testing.—The anti-doping enforcement agency shall perform and manage test distribution planning (including intelligence-based testing), the sample collection process, and incompetition and out-of-competition testing (including no-advance-notice testing).

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1 (D) Testing Laboratories.—The anti-2 doping and medication control enforcement 3 agency shall accredit testing laboratories based 4 upon the standards established under this Act, 5 and shall monitor, test, and audit accredited 6 laboratories to ensure continuing compliance 7 with accreditation standards. 8 (5) Anti-doping and medication control 9 STANDING COMMITTEE.—The anti-doping and medi-10 cation control standing committee shall, in consulta-11 tion with the anti-doping and medication control en-12 forcement agency, develop lists of permitted and pro-13 hibited medications, methods, and substances for 14 recommendation to, and approval by, the Authority. 15 Any such list may prohibit the administration of any 16 substance or method to a horse at any time after 17 such horse becomes a covered horse if the Authority 18 determines such substance or method has a long-19 term degrading effect on the soundness of a horse. 20 (d) Prohibition.—Except as provided in sub-21 sections (e) and (f), the horseracing anti-doping and medi-22 cation control program shall prohibit the administration 23 of any prohibited or otherwise permitted substance to a covered horse within 48 hours of its next racing start, ef-

fective as of the program effective date.

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1 (e) Advisory Committee Study and Report.— 2 (1) In General.—Not later than the program 3 effective date, the Authority shall convene an advi-4 sory committee comprised of horseracing anti-doping 5 and medication control industry experts, including a 6 member designated by the anti-doping and medica-7 tion control enforcement agency, to conduct a study 8 on the use of furosemide on horses during the 48-9 hour period before the start of a race, including the 10 effect of furosemide on equine health and the integ-11 rity of competition and any other matter the Author-12 ity considers appropriate. 13 (2) Report.—Not later than three years after 14 the program effective date, the Authority shall direct 15 the advisory committee convened under paragraph 16 (1) to submit to the Authority a written report on 17 the study conducted under that paragraph that in-18 cludes recommended changes, if any, to the prohibi-19 tion in subsection (d). 20 (3) Modification of Prohibition.— 21 (A) IN GENERAL.—After receipt of the re-22 port required by paragraph (2), the Authority 23 may, by unanimous vote of the Board of the 24 Authority, modify the prohibition in subsection 25 (d) and, notwithstanding subsection (f), any Case 1:20-cr-00160-MKV Document 327-7 Filed 02/05/21 Page 40 of 65 SEN. APPRO.

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1	such modification shall apply to all States be-
2	ginning on the date that is three years after the
3	program effective date.
4	(B) Condition.—In order for a unani-
5	mous vote described in subparagraph (A) to ef-
6	fect a modification of the prohibition in sub-
7	section (d), the vote must include unanimous
8	adoption of each of the following findings:
9	(i) That the modification is war-
10	ranted.
11	(ii) That the modification is in the
12	best interests of horse racing.
13	(iii) That furosemide has no perform-
14	ance enhancing effect on individual horses.
15	(iv) That public confidence in the in-
16	tegrity and safety of racing would not be
17	adversely affected by the modification.
18	(f) Exemption.—
19	(1) In general.—Except as provided in para-
20	graph (2), only during the three-year period begin-
21	ning on the program effective date, a State racing
22	commission may submit to the Authority, at such
23	time and in such manner as the Authority may re-
24	quire, a request for an exemption from the prohibi-

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1	tion in subsection (d) with respect to the use of
2	furosemide on covered horses during such period.
3	(2) Exceptions.—An exemption under para-
4	graph (1) may not be requested for—
5	(A) two-year-old covered horses; or
6	(B) covered horses competing in stakes
7	races.
8	(3) Contents of request.—A request under
9	paragraph (1) shall specify the applicable State rac-
10	ing commission's requested limitations on the use of
11	furosemide that would apply to the State under the
12	horseracing anti-doping and medication control pro-
13	gram during such period. Such limitations shall be
14	no less restrictive on the use and administration of
15	furosemide than the restrictions set forth in State's
16	laws and regulations in effect as of September 1,
17	2020.
18	(4) Grant of exemption.—Subject to sub-
19	section (e)(3), the Authority shall grant an exemp-
20	tion requested under paragraph (1) for the remain-
21	der of such period and shall allow the use of
22	furosemide on covered horses in the applicable State,
23	in accordance with the requested limitations.
24	(g) Baseline Anti-doping and Medication Con-
25	TROL RILLES.—

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1	(1) In General.—Subject to paragraph (3),
2	the baseline anti-doping and medication control rules
3	described in paragraph (2) shall—
4	(A) constitute the initial rules of the horse-
5	racing anti-doping and medication control pro-
6	gram; and
7	(B) except as exempted pursuant to sub-
8	sections (e) and (f), remain in effect at all
9	times after the program effective date.
10	(2) Baseline anti-doping medication con-
11	TROL RULES DESCRIBED.—
12	(A) In General.—The baseline anti-
13	doping and medication control rules described
14	in this paragraph are the following:
15	(i) The lists of permitted and prohib-
16	ited substances (including drugs, medica-
17	tions, and naturally occurring substances
18	and synthetically occurring substances) in
19	effect for the International Federation of
20	Horseracing Authorities, including the
21	International Federation of Horseracing
22	Authorities International Screening Limits
23	for urine, dated May 2019, and the Inter-
24	national Federation of Horseracing Au-

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1	thorities International Screening Limits for
2	plasma, dated May 2019.
3	(ii) The World Anti-Doping Agency
4	International Standard for Laboratories
5	(version 10.0), dated November 12, 2019.
6	(iii) The Association of Racing Com-
7	missioners International out-of-competition
8	testing standards, Model Rules of Racing
9	(version 9.2).
10	(iv) The Association of Racing Com-
11	missioners International penalty and mul-
12	tiple medication violation rules, Model
13	Rules of Racing (version 6.2).
14	(B) CONFLICT OF RULES.—In the case of
15	a conflict among the rules described in subpara-
16	graph (A), the most stringent rule shall apply.
17	(3) Modifications to baseline rules.—
18	(A) DEVELOPMENT BY ANTI-DOPING AND
19	MEDICATION CONTROL STANDING COM-
20	MITTEE.—The anti-doping and medication con-
21	trol standing committee, in consultation with
22	the anti-doping and medication control enforce-
23	ment agency, may develop and submit to the
24	Authority for approval by the Authority pro-

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1	posed modifications to the baseline anti-doping
2	and medication control rules.
3	(B) AUTHORITY APPROVAL.—If the Au-
4	thority approves a proposed modification under
5	this paragraph, the proposed modification shall
6	be submitted to and considered by the Commis-
7	sion in accordance with section 1204.
8	(C) Anti-doping and medication con-
9	TROL ENFORCEMENT AGENCY VETO AUTHOR-
10	ITY.—The Authority shall not approve any pro-
11	posed modification that renders an anti-doping
12	and medication control rule less stringent than
13	the baseline anti-doping and medication control
14	rules described in paragraph (2) (including by
15	increasing permitted medication thresholds,
16	adding permitted medications, removing prohib-
17	ited medications, or weakening enforcement
18	mechanisms) without the approval of the anti-
19	doping and medication control enforcement
20	agency.
21	SEC. 1207. RACETRACK SAFETY PROGRAM.
22	(a) Establishment and Considerations.—
23	(1) In general.—Not later than the program
24	effective date, and after notice and an opportunity
25	for public comment in accordance with section 1204,

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1	the Authority shall establish a racetrack safety pro-
2	gram applicable to all covered horses, covered per-
3	sons, and covered horseraces in accordance with the
4	registration of covered persons under section
5	1205(d).
6	(2) Considerations in development of
7	SAFETY PROGRAM.—In the development of the
8	horseracing safety program for covered horses, cov-
9	ered persons, and covered horseraces, the Authority
10	and the Commission shall take into consideration ex-
11	isting safety standards including the National Thor-
12	oughbred Racing Association Safety and Integrity
13	Alliance Code of Standards, the International Fed-
14	eration of Horseracing Authority's International
15	Agreement on Breeding, Racing, and Wagering, and
16	the British Horseracing Authority's Equine Health
17	and Welfare program.
18	(b) Elements of Horseracing Safety Pro-
19	GRAM.—The horseracing safety program shall include the
20	following:
21	(1) A set of training and racing safety stand-
22	ards and protocols taking into account regional dif-
23	ferences and the character of differing racing facili-
24	ties.

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1	(2) A uniform set of training and racing safety
2	standards and protocols consistent with the humane
3	treatment of covered horses, which may include lists
4	of permitted and prohibited practices or methods
5	(such as crop use).
6	(3) A racing surface quality maintenance sys-
7	tem that—
8	(A) takes into account regional differences
9	and the character of differing racing facilities;
10	and
11	(B) may include requirements for track
12	surface design and consistency and established
13	standard operating procedures related to track
14	surface, monitoring, and maintenance (such as
15	standardized seasonal assessment, daily track-
16	ing, and measurement).
17	(4) A uniform set of track safety standards and
18	protocols, that may include rules governing oversight
19	and movement of covered horses and human and
20	equine injury reporting and prevention.
21	(5) Programs for injury and fatality data anal-
22	ysis, that may include pre- and post-training and
23	race inspections, use of a veterinarian's list, and
24	concussion protocols.

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1	(6) The undertaking of investigations at race-
2	track and non-racetrack facilities related to safety
3	violations.
4	(7) Procedures for investigating, charging, and
5	adjudicating violations and for the enforcement of
6	civil sanctions for violations.
7	(8) A schedule of civil sanctions for violations.
8	(9) Disciplinary hearings, which may include
9	binding arbitration, civil sanctions, and research.
10	(10) Management of violation results.
11	(11) Programs relating to safety and perform-
12	ance research and education.
13	(12) An evaluation and accreditation program
14	that ensures that racetracks in the United States
15	meet the standards described in the elements of the
16	Horseracing Safety Program.
17	(c) Activities.—The following activities shall be car-
18	ried out under the racetrack safety program:
19	(1) Standards for racetrack safety.—
20	The development, by the racetrack safety standing
21	committee of the Authority in section 1203(c)(2) of
22	uniform standards for racetrack and horseracing
23	safety.
24	(2) Standards for safety and perform-
25	ANCE ACCREDITATION.—

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1	(A) IN GENERAL.—Not later than 120
2	days before the program effective date, the Au-
3	thority, in consultation with the racetrack safe-
4	ty standing committee, shall issue, by rule in
5	accordance with section 1204—
6	(i) safety and performance standards
7	of accreditation for racetracks; and
8	(ii) the process by which a racetrack
9	may achieve and maintain accreditation by
10	the Authority.
11	(B) Modifications.—
12	(i) IN GENERAL.—The Authority may
13	modify rules establishing the standards
14	issued under subparagraph (A), as the Au-
15	thority considers appropriate.
16	(ii) NOTICE AND COMMENT.—The
17	Commission shall publish in the Federal
18	Register any proposed rule of the Author-
19	ity, and provide an opportunity for public
20	comment with respect to, any modification
21	under clause (i) in accordance with section
22	1204.
23	(C) EXTENSION OF PROVISIONAL OR IN-
24	TERIM ACCREDITATION.—The Authority may,
25	by rule in accordance with section 1204, extend

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1	provisional or interim accreditation to a race-
2	track accredited by the National Thoroughbred
3	Racing Association Safety and Integrity Alli-
4	ance on a date before the program effective
5	date.
6	(3) Nationwide safety and performance
7	DATABASE.—
8	(A) IN GENERAL.—Not later than one year
9	after the program effective date, and after no-
10	tice and an opportunity for public comment in
11	accordance with section 1204, the Authority, in
12	consultation with the Commission, shall develop
13	and maintain a nationwide database of race-
14	horse safety, performance, health, and injury
15	information for the purpose of conducting an
16	epidemiological study.
17	(B) Collection of Information.—In
18	accordance with the registration of covered per-
19	sons under section 1205(d), the Authority may
20	require covered persons to collect and submit to
21	the database described in subparagraph (A)
22	such information as the Authority may require
23	to further the goal of increased racehorse wel-
24	fare.

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1	SEC. 1208. RULE VIOLATIONS AND CIVIL SANCTIONS.
2	(a) Description of Rule Violations.—
3	(1) In general.—The Authority shall issue, by
4	rule in accordance with section 1204, a description
5	of safety, performance, and anti-doping and medica-
6	tion control rule violations applicable to covered
7	horses and covered persons.
8	(2) Elements.—The description of rule viola-
9	tions established under paragraph (1) may include
10	the following:
11	(A) With respect to a covered horse, strict
12	liability for covered trainers for—
13	(i) the presence of a prohibited sub-
14	stance or method in a sample or the use of
15	a prohibited substance or method;
16	(ii) the presence of a permitted sub-
17	stance in a sample in excess of the amount
18	allowed by the horseracing anti-doping and
19	medication control program; and
20	(iii) the use of a permitted method in
21	violation of the applicable limitations es-
22	tablished under the horseracing anti-
23	doping and medication control program.
24	(B) Attempted use of a prohibited sub-
25	stance or method on a covered horse.

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1	(C) Possession of any prohibited substance
2	or method.
3	(D) Attempted possession of any prohib-
4	ited substance or method.
5	(E) Administration or attempted adminis-
6	tration of any prohibited substance or method
7	on a covered horse.
8	(F) Refusal or failure, without compelling
9	justification, to submit a covered horse for sam-
10	ple collection.
11	(G) Failure to cooperate with the Author-
12	ity or an agent of the Authority during any in-
13	vestigation.
14	(H) Failure to respond truthfully, to the
15	best of a covered person's knowledge, to a ques-
16	tion of the Authority or an agent of the Author-
17	ity with respect to any matter under the juris-
18	diction of the Authority.
19	(I) Tampering or attempted tampering
20	with the application of the safety, performance,
21	or anti-doping and medication control rules or
22	process adopted by the Authority, including—
23	(i) the intentional interference, or an
24	attempt to interfere, with an official or
25	agent of the Authority;

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1	(ii) the procurement or the provision
2	of fraudulent information to the Authority
3	or agent; and
4	(iii) the intimidation of, or an attempt
5	to intimidate, a potential witness.
6	(J) Trafficking or attempted trafficking in
7	any prohibited substance or method.
8	(K) Assisting, encouraging, aiding, abet-
9	ting, conspiring, covering up, or any other type
10	of intentional complicity involving a safety, per-
11	formance, or anti-doping and medication control
12	rule violation or the violation of a period of sus-
13	pension or eligibility.
14	(L) Threatening or seeking to intimidate a
15	person with the intent of discouraging the per-
16	son from the good faith reporting to the Au-
17	thority, an agent of the Authority or the Com-
18	mission, or the anti-doping and medication con-
19	trol enforcement agency under section 1205(e),
20	of information that relates to—
21	(i) an alleged safety, performance, or
22	anti-doping and medication control rule
23	violation; or

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1	(ii) alleged noncompliance with a safe-	
2	ty, performance, or anti-doping and medi-	
3	cation control rule.	
4	(b) Testing Laboratories.—	
5	(1) Accreditation and standards.—Not	
6	later than 120 days before the program effective	
7	date, the Authority shall, in consultation with the	
8	anti-doping and medication control enforcement	
9	agency, establish, by rule in accordance with section	
10	1204—	
11	(A) standards of accreditation for labora-	
12	tories involved in testing samples from covered	
13	horses;	
14	(B) the process for achieving and main-	
15	taining accreditation; and	
16	(C) the standards and protocols for testing	
17	such samples.	
18	(2) Administration.—The accreditation of	
19	laboratories and the conduct of audits of accredited	
20	laboratories to ensure compliance with Authority	
21	rules shall be administered by the anti-doping and	
22	medication control enforcement agency. The anti-	
23	doping and medication control enforcement agency	
24	shall have the authority to require specific test sam-	

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1	ples to be directed to and tested by laboratories hav-
2	ing special expertise in the required tests.
3	(3) Extension of provisional or interim
4	ACCREDITATION.—The Authority may, by rule in ac-
5	cordance with section 1204, extend provisional or in-
6	terim accreditation to a laboratory accredited by the
7	Racing Medication and Testing Consortium, Inc., on
8	a date before the program effective date.
9	(4) Selection of Laboratories.—
10	(A) In general.—Except as provided in
11	paragraph (2), a State racing commission may
12	select a laboratory accredited in accordance
13	with the standards established under paragraph
14	(1) to test samples taken in the applicable
15	State.
16	(B) Selection by the authority.—If a
17	State racing commission does not select an ac-
18	credited laboratory under subparagraph (A),
19	the Authority shall select such a laboratory to
20	test samples taken in the State concerned.
21	(c) Results Management and Disciplinary
22	Process.—
23	(1) In general.—Not later than 120 days be-
24	fore the program effective date, the Authority shall
25	establish in accordance with section 1204—

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1	(A) rules for safety, performance, and anti-
2	doping and medication control results manage-
3	ment; and
4	(B) the disciplinary process for safety, per-
5	formance, and anti-doping and medication con-
6	trol rule violations.
7	(2) Elements.—The rules and process estab-
8	lished under paragraph (1) shall include the fol-
9	lowing:
10	(A) Provisions for notification of safety,
11	performance, and anti-doping and medication
12	control rule violations.
13	(B) Hearing procedures.
14	(C) Standards for burden of proof.
15	(D) Presumptions.
16	(E) Evidentiary rules.
17	(F) Appeals.
18	(G) Guidelines for confidentiality and pub-
19	lic reporting of decisions.
20	(3) Due process.—The rules established
21	under paragraph (1) shall provide for adequate due
22	process, including impartial hearing officers or tribu-
23	nals commensurate with the seriousness of the al-
24	leged safety, performance, or anti-doping and medi-

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1	cation control rule violation and the possible civil
2	sanctions for such violation.
3	(d) CIVIL SANCTIONS.—
4	(1) In general.—The Authority shall estab-
5	lish uniform rules, in accordance with section 1204,
6	imposing civil sanctions against covered persons or
7	covered horses for safety, performance, and anti-
8	doping and medication control rule violations.
9	(2) Requirements.—The rules established
10	under paragraph (1) shall—
11	(A) take into account the unique aspects of
12	horseracing;
13	(B) be designed to ensure fair and trans-
14	parent horseraces; and
15	(C) deter safety, performance, and anti-
16	doping and medication control rule violations.
17	(3) Severity.—The civil sanctions under para-
18	graph (1) may include—
19	(A) lifetime bans from horseracing,
20	disgorgement of purses, monetary fines and
21	penalties, and changes to the order of finish in
22	covered races; and
23	(B) with respect to anti-doping and medi-
24	cation control rule violators, an opportunity to
25	reduce the applicable civil sanctions that is

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1	comparable to the opportunity provided by the
2	Protocol for Olympic Movement Testing of the
3	United States Anti-Doping Agency.
4	(e) Modifications.—The Authority may propose a
5	modification to any rule established under this section as
6	the Authority considers appropriate, and the proposed
7	modification shall be submitted to and considered by the
8	Commission in accordance with section 1204.
9	SEC. 1209. REVIEW OF FINAL DECISIONS OF THE AUTHOR-
10	ITY.
11	(a) Notice of Civil Sanctions.— If the Authority
12	imposes a final civil sanction for a violation committed by
13	a covered person pursuant to the rules or standards of
14	the Authority, the Authority shall promptly submit to the
15	Commission notice of the civil sanction in such form as
16	the Commission may require.
17	(b) REVIEW BY ADMINISTRATIVE LAW JUDGE.—
18	(1) In general.—With respect to a final civil
19	sanction imposed by the Authority, on application by
20	the Commission or a person aggrieved by the civil
21	sanction filed not later than 30 days after the date
22	on which notice under subsection (a) is submitted,
23	the civil sanction shall be subject to de novo review
24	by an administrative law judge.
25	(2) Nature of Review.—

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1	(A) In General.—In matters reviewed
2	under this subsection, the administrative law
3	judge shall determine whether—
4	(i) a person has engaged in such acts
5	or practices, or has omitted such acts or
6	practices, as the Authority has found the
7	person to have engaged in or omitted;
8	(ii) such acts, practices, or omissions
9	are in violation of this Act or the anti-
10	doping and medication control or racetrack
11	safety rules approved by the Commission;
12	or
13	(iii) the final civil sanction of the Au-
14	thority was arbitrary, capricious, an abuse
15	of discretion, or otherwise not in accord-
16	ance with law.
17	(B) Conduct of Hearing.—An adminis-
18	trative law judge shall conduct a hearing under
19	this subsection in such a manner as the Com-
20	mission may specify by rule, which shall con-
21	form to section 556 of title 5, United States
22	Code.
23	(3) Decision by administrative law
24	JUDGE.—

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1	(A) IN GENERAL.—With respect to a mat-
2	ter reviewed under this subsection, an adminis-
3	trative law judge—
4	(i) shall render a decision not later
5	than 60 days after the conclusion of the
6	hearing;
7	(ii) may affirm, reverse, modify, set
8	aside, or remand for further proceedings,
9	in whole or in part, the final civil sanction
10	of the Authority; and
11	(iii) may make any finding or conclu-
12	sion that, in the judgment of the adminis-
13	trative law judge, is proper and based on
14	the record.
15	(B) FINAL DECISION.—A decision under
16	this paragraph shall constitute the decision of
17	the Commission without further proceedings
18	unless a notice or an application for review is
19	timely filed under subsection (c).
20	(c) Review by Commission.—
21	(1) Notice of review by commission.—The
22	Commission may, on its own motion, review any de-
23	cision of an administrative law judge issued under
24	subsection (b)(3) by providing written notice to the
25	Authority and any interested party not later than 30

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1	days after the date on which the administrative law
2	judge issues the decision.
3	(2) Application for review.—
4	(A) In general.—The Authority or a per-
5	son aggrieved by a decision issued under sub-
6	section (b)(3) may petition the Commission for
7	review of such decision by filing an application
8	for review not later than 30 days after the date
9	on which the administrative law judge issues
10	the decision.
11	(B) EFFECT OF DENIAL OF APPLICATION
12	FOR REVIEW.—If an application for review
13	under subparagraph (A) is denied, the decision
14	of the administrative law judge shall constitute
15	the decision of the Commission without further
16	proceedings.
17	(C) Discretion of commission.—
18	(i) In general.—A decision with re-
19	spect to whether to grant an application
20	for review under subparagraph (A) is sub-
21	ject to the discretion of the Commission.
22	(ii) Matters to be considered.—
23	In determining whether to grant such an
24	application for review, the Commission

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1	shall consider whether the application
2	makes a reasonable showing that—
3	(I) a prejudicial error was com-
4	mitted in the conduct of the pro-
5	ceeding; or
6	(II) the decision involved—
7	(aa) an erroneous applica-
8	tion of the anti-doping and medi-
9	cation control or racetrack safety
10	rules approved by the Commis-
11	sion; or
12	(bb) an exercise of discretion
13	or a decision of law or policy that
14	warrants review by the Commis-
15	sion.
16	(3) Nature of Review.—
17	(A) In General.—In matters reviewed
18	under this subsection, the Commission may—
19	(i) affirm, reverse, modify, set aside,
20	or remand for further proceedings, in
21	whole or in part, the decision of the admin-
22	istrative law judge; and
23	(ii) make any finding or conclusion
24	that, in the judgement of the Commission,
25	is proper and based on the record.

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1	(B) DE NOVO REVIEW.—The Commission
2	shall review de novo the factual findings and
3	conclusions of law made by the administrative
4	law judge.
5	(C) Consideration of additional evi-
6	DENCE.—
7	(i) MOTION BY COMMISSION.—The
8	Commission may, on its own motion, allow
9	the consideration of additional evidence.
10	(ii) MOTION BY A PARTY.—
11	(I) In general.—A party may
12	file a motion to consider additional
13	evidence at any time before the
14	issuance of a decision by the Commis-
15	sion, which shall show, with particu-
16	larity, that—
17	(aa) such additional evidence
18	is material; and
19	(bb) there were reasonable
20	grounds for failure to submit the
21	evidence previously.
22	(II) Procedure.—The Commis-
23	sion may—
24	(aa) accept or hear addi-
25	tional evidence; or

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1	(bb) remand the proceeding
2	to the administrative law judge
3	for the consideration of addi-
4	tional evidence.
5	(d) STAY OF PROCEEDINGS.—Review by an adminis-
6	trative law judge or the Commission under this section
7	shall not operate as a stay of a final civil sanction of the
8	Authority unless the administrative law judge or Commis-
9	sion orders such a stay.
10	SEC. 1210. UNFAIR OR DECEPTIVE ACTS OR PRACTICES.
11	The sale of a covered horse, or of any other horse
12	in anticipation of its future participation in a covered race,
13	shall be considered an unfair or deceptive act or practice
14	in or affecting commerce under section 5(a) of the Federal
15	Trade Commission Act (15 U.S.C. 45(a)) if the seller—
16	(1) knows or has reason to know the horse has
17	been administered—
18	(A) a bisphosphonate prior to the horse's
19	fourth birthday; or
20	(B) any other substance or method the Au-
21	thority determines has a long-term degrading
22	effect on the soundness of the covered horse;
23	and

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1	(2) fails to disclose to the buyer the administra-
2	tion of the bisphosphonate or other substance or
3	method described in paragraph (1)(B).
4	SEC. 1211. STATE DELEGATION; COOPERATION.
5	(a) State Delegation.—
6	(1) In General.—The Authority may enter
7	into an agreement with a State racing commission to
8	implement, within the jurisdiction of the State rac-
9	ing commission, a component of the racetrack safety
10	program or, with the concurrence of the anti-doping
11	and medication control enforcement agency under
12	section 1205(e), a component of the horseracing
13	anti-doping and medication control program, if the
14	Authority determines that the State racing commis-
15	sion has the ability to implement such component in
16	accordance with the rules, standards, and require-
17	ments established by the Authority.
18	(2) Implementation by state racing com-
19	MISSION.—A State racing commission or other ap-
20	propriate regulatory body of a State may not imple-
21	ment such a component in a manner less restrictive
22	than the rule, standard, or requirement established
23	by the Authority.
24	(b) Cooperation.—To avoid duplication of func-
25	tions, facilities, and personnel, and to attain closer coordi-

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- 2 tration of Federal and State law, where conduct by any
- 3 person subject to the horseracing medication control pro-
- 4 gram or the racetrack safety program may involve both
- 5 a medication control or racetrack safety rule violation and
- 6 violation of Federal or State law, the Authority and Fed-
- 7 eral or State law enforcement authorities shall cooperate
- 8 and share information.

9 SEC. 1212. DETERMINATION OF BUDGETARY EFFECTS.

- The budgetary effects of this Act, for the purpose of
- 11 complying with the Statutory Pay-As-You-Go Act of 2010,
- 12 shall be determined by reference to the latest statement
- 13 titled "Budgetary Effects of PAYGO Legislation" for this
- 14 Act, submitted for printing in the Congressional Record
- 15 by the Chairman of the House Budget Committee, pro-
- 16 vided that such statement has been submitted prior to the
- 17 vote on passage.

18 TITLE XIII—COMMUNITY

19 **DEVELOPMENT BLOCK GRANTS**

- 20 SEC. 1301. COMMUNITY DEVELOPMENT BLOCK GRANTS.
- 21 (a) IN GENERAL.—Funds previously made available
- 22 in chapter 9 of title X of the Disaster Relief Appropria-
- 23 tions Act, 2013 (Public Law 113-2, division A; 127 Stat.
- 24 36) under the heading "DEPARTMENT OF HOUSING
- 25 AND URBAN DEVELOPMENT—Community Planning